IN THE CLAIMS:

Please amend claims 1 and 34 as follows:

- 1. (Currently Amended) A compound comprising two or more antigen binding regions linked to at least one prodrug-activating enzyme, wherein
 - a) the antigen binding regions consist of a single polypeptide chain;
 - the single polypeptide chain is comprised of a first variable domain, a second variable domain, and a polypeptide linker connecting the first variable domain and the second variable domain; wherein a nucleotide sequence encoding the polypeptide linker is formed by two partially overlapping PCR primers during a PCR reaction that links the first variable domain and the second variable domain; and wherein
 - c) a nucleotide sequence encoding the polypeptide linker is formed by
 two partially overlapping PCR primers during a PCR reaction that
 links the first variable domain and the second variable domain; and
 wherein
 - d) c) the compound has a bivalent or a multivalent structure.
- 2. (Previously Presented) A compound as claimed in claim 1, wherein the compound further comprises covalently bonded carbohydrates.
- 3. (Previously Presented) A compound as claimed in claim 1, wherein at least one antigen binding region comprises a variable domain of a heavy antibody chain and a variable domain of a light antibody chain (sFv fragment).

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1300 L Street NW Washington, DC 20005 202,408,4000 Fax 202,408,4400 www.finnegan.com 4. (Original) A compound as claimed in claim 1, wherein the antigen binding region binds to a tumor-associated antigen (TAA).

5. (Previously Presented) A compound as claimed in claim 4, wherein the TAA is selected from the group consisting of an N-CAM, PEM, EGF-R, Sialyl-Le^a, Sialyl-Le^x, TFβ, GICA, GD₃, GD₂, TAG72, CA125, the 24-25 kDa glycoprotein defined by Mab L6, and CEA.

6. (Previously Presented) A compound as claimed in claim 1, wherein the enzyme is selected from the group consisting of a lactamase, pyroglutamate aminopeptidase, D-aminopeptidase, oxidase, peroxidase, phosphatase, hydroxynitrile lyase, protease, esterase, carboxypeptidase and glycosidase.

7. (Previously Presented) A compound as claimed in claim 6, wherein the enzyme is a β -glucuronidase, which is selected from the group consisting of an E. coli β -glucuronidase, a Kobayasia nipponica β -glucuronidase, a Secale Cereale β -glucuronidase and a human β -glucuronidase.

- 8. (Original) A compound as claimed in claim 1, wherein the antigen binding region is linked to the enzyme via a peptide linker.
- 9. (Previously Presented) A compound as claimed in claim 2, wherein glycosylation covalently bonds the carbohydrates to the compound, and the glycosylation takes place either by means of chemical methods or by a selection of suitable expression systems.

10. (Previously Presented) A compound as claimed in claim 1, which has undergone secretory expression in *Saccharomyces cerevisiae* or in *Hansenula polymorpha*.

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- 11. (Previously Presented) A compound as claimed in claim 1, which is expressed in *E. coli* and is subsequently chemically glycosylated.
- 12. (Previously Presented) A compound as claimed in claim 30, wherein the $sFv-\beta$ -lactamase fusion protein has undergone periplasmic expression in *E. coli* and is chemically glycosylated.
- 13. (Previously Presented) A compound as claimed in claim 30, wherein the sFv-β-lactamase fusion protein has undergone secretory expression in *Saccharomyces* cerevisiae or *Hansenula polymorpha*.
 - 14. (Withdrawn) A nucleic acid coding for a compound as claimed in claim 1.
- 15. (Withdrawn) A nucleic acid as claimed in claim 14, coding for a humanized sFv fragment against CEA and a human β -glucuronidase.
- 16. (Withdrawn) A nucleic acid as claimed in claim 14 with the sequence CCAAGCTTAT GAATATGCAA ATCCTGCTCA TGAATATGCA AATCCTCTGA 50 ATCTACATGG TAAATATAGG TTTGTCTATA CCACAAACAG AAAAACATGA 100 GATCACAGTT CTCTCTACAG TTACTGAGCA CACAGGACCT CACC ATG GGA TGG 153 Met Glv Trp AGC TGT ATC ATC CTC TTC TTG GTA GCA ACA GCT ACA GGTAAGGGGC 199 Ser Cys lle lle Leu Phe Leu Val Ala Thr Ala Thr -10 TCACAGTAGC AGGCTTGAGG TCTGGACATA TATATGGGTG ACAATAGACAT 249 CCACTTTGCC TTTCTCCA CA GGT GTC CAC TCC CAG GTC CAA CTG CAG 298 Gly Val His Ser Gln Val Gln Leu Gln GAG AGC GGT CCA GGT CTT GTG AGA CCT AGC CAG ACC CTG AGC CTG 343 Glu Ser Gly Pro Gly Leu Val Arg Pro Ser Gln Thr ACC TGC ACC GTG TCT GGC TTC ACC ATC AGC AGT GGT TAT AGC TGG 388 Thr Cys Thr Val Ser Gly Phe Thr Ile Ser Ser Gly Tyr Ser Trp CAC TGG GTG AGA CAG CCA CCT GGA CGA GGT CTT GAG TGG ATT GGA 433 Val Arg Gln Pro Pro Gly Arg Gly Leu Glu Trp Ile His Trp 40

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	TAC Tyr								AAC ⁻ Asn								478
									ACC Thr	AGC						r	523
					AGC				C GCC Ala						T T	AT	568
									CAC His						TGC Trp 110		613
									C TCC I Sei		r <u>Gly</u>						658
						r Gly			GC GC y GI								703
1	Gln	Ser	Pro	Ser	Ser	Leu	Ser	Ala	C AG Ser	Val 150	Gly O	Asp	Arg	l Va	ı T	hr hr	748
	ATC lle								GTA Val								793
									A AAG Lys		Le						838
			CTG Leu			Gly			AGC Ser							у	883
									ATC ile								928
	GAC Asp					Tyr			CAG Gln							r	973
									G ATC lle		GGT	GAG	TAGA	`TA	TTA	AACTTT	1023
•	TGC	TTCC	TCA	GTT	GGA1	CTG	AGT	AACT	rccc	AAT	CTTC	TCT	CTG(CTC AAA Leu Lys	1077
								His	ACA Thr								1119
GGTAAGCCAG CCCAGGACTC GCCCTCCAGC TCAAGGCGGG ACAAGAGCCC										1169							
-	TAG	AGTO	GCC	TGA	GTC	CAG	G GA	CAG	GCCC	AGC	AGG	GTG	C TG	ACG	CAT	CC	1219

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ACCTCCATCC CAGATCCCCG TAACTCCCAA TCTTCTCTCT GCA GCG GCG GCG 1 Ala Ala Ala 260	1271
	1316
GAG TGC AAG GAG CTG GAC GGC CTC TGG AGC TTC CGC GCC GAC TTC Glu Cys Lys Glu Leu Asp Gly Leu Trp Ser Phe Arg Ala Asp Phe 280 290	1361
Ser Asp Asn Arg Arg Gly Phe Glu Glu Gln Trp Tyr Arg Arg 300	1406
CCG CTG TGG GAG TCA GGC CCC ACC GTG GAC ATG CCA GTT CCC TCC Pro Leu Trp Glu Ser Gly Pro Thr Val Asp Met Pro Val Pro Ser 310 320	1451
AGC TTC AAT GAC ATC AGC CAG GAC TGG CGT CTG CGG CAT TTT GTC Ser Phe Asn Asp IIe Ser Gln Asp Trp Arg Leu Arg His Phe Val 330	1496
GGC TGG GTG TGG TAC GAA CGG GAG GTG ATC CTG CCG GAG CGA TGG Gly Trp Val Trp Tyr Glu Arg Glu Val IIe Leu Pro Glu Arg Trp 340 350	1541
ACC CAG GAC CTG CGC ACA AGA GTG GTG CTG AGG ATT GGC AGT GCC Thr Gin Asp Leu Arg Thr Arg Val Val Leu Arg Ile Gly Ser Ala 360	1586
CAT TCC TAT GCC ATC GTG TGG GTG AAT GGG GTC GAC ACG CTA GAG His Ser Tyr Ala IIe Val Trp Val Asn Gly Val Asp Thr Leu Glu 370 380	1631
CAT GAG GGG GGC TAC CTC CCC TTC GAG GCC GAC ATC AGC AAC CTG His Glu Gly Gly Tyr Leu Pro Phe Glu Ala Asp Ile Ser Asn Leu 390	1676
GTC CAG GTG GGG CCC CTG CCC TCC CGG CTC CGA ATC ACT ATC GCC Val Gln Val Gly Pro Leu Pro Ser Arg Leu Arg Ile Thr Ile Ala 400 410	1721
	1766
	1811
	1856
	1901
	1946
	1991

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CTT TTG GAT GCA GAA AAC AAA GTC GTG GCG AAT GGG ACT GGG ACC Leu Leu Asp Ala Glu Asn Lys Val Val Ala Asn Gly Thr Gly Thr 510	2036
CAG GGC CAA CTT AAG GTG CCA GGT GTC AGC CTC TGG TGG CCG TAC GIn Gly Gln Leu Lys Val Pro Gly Val Ser Leu Trp Trp Pro Tyr 520 530	2081
CTG ATG CAC GAA CGC CCT GCC TAT CTG TAT TCA TTG GAG GTG CAG Leu Met His Glu Arg Pro Ala Tyr Leu Tyr Ser Leu Glu Val Gln 540	2126
CTG ACT GCA CAG ACG TCA CTG GGG CCT GTG TCT GAC TTC TAC ACA Leu Thr Ala Gln Thr Ser Leu Gly Pro Val Ser Asp Phe Tyr Thr 550 560	2171
CTC CCT GTG GGG ATC CGC ACT GTG GCT GTC ACC AAG AGC CAG TTC Leu Pro Val Gly lle Arg Thr Val Ala Val Thr Lys Ser Gln Phe 570	2216
CTC ATC AAT GGG AAA CCT TTC TAT TTC CAC GGT GTC AAC AAG CAT Leu lle Asn Gly Lys Pro Phe Tyr Phe His Gly Val Asn Lys His 580	2261
GAG GAT GCG GAC ATC CGA GGG AAG GGC TTC GAC TGG CCG CTG CTG Glu Asp Ala Asp Ile Arg Gly Lys Gly Phe Asp Trp Pro Leu Leu 600	2306
GTG AAG GAC TTC AAC CTG CTT CGC TGG CTT GGT GCC AAC GCT TTC Val Lys Asp Phe Asn Leu Leu Arg Trp Leu Gly Ala Asn Ala Phe 610 620	2351
CGT ACC AGC CAC TAC CCC TAT GCA GAG GAA GTG ATG CAG ATG TGT Arg Thr Ser His Tyr Pro Tyr Ala Glu Glu Val Met Gln Met Cys 630	2396
GAC CGC TAT GGG ATT GTG GTC ATC GAT GAG TGT CCC GGC GTG GGC Asp Arg Tyr Gly lle Val Val lle Asp Glu Cys Pro Gly Val Gly 640 650	2441
CTG GCG CTG CCG CAG TTC TTC AAC AAC GTT TCT CTG CAT CAC CAC Leu Ala Leu Pro Gln Phe Phe Asn Asn Val Ser Leu His His His 660	2486
ATG CAG GTG ATG GAA GAA GTG GTG CGT AGG GAC AAG AAC CAC CCC Met Gln Val Met Glu Glu Val Val Arg Arg Asp Lys Asn His Pro 670 680	2531
GCG GTC GTG ATG TGG TCT GTG GCC AAC GAG CCT GCG TCC CAC CTA Ala Val Val Met Trp Ser Val Ala Asn Glu Pro Ala Ser His Leu 690	2576
GAA TCT GCT GGC TAC TAC TTG AAG ATG GTG ATC GCT CAC ACC AAA Glu Ser Ala Gly Tyr Tyr Leu Lys Met Val Ile Ala His Thr Lys 700 710	2621
TCC TTG GAC CCC TCC CGG CCT GTG ACC TTT GTG AGC AAC TCT AAC Ser Leu Asp Pro Ser Arg Pro Val Thr Phe Val Ser Asn Ser Asn 720	2666
TAT GCA GCA GAC AAG GGG GCT CCG TAT GTG GAT GTG ATC TGT TTG Tyr Ala Ala Asp Lys Gly Ala Pro Tyr Val Asp Val Ile Cys Leu 730 740	2711
AAC AGC TAC TAC TCT TGG TAT CAC GAC TAC GGG CAC CTG GAG TTG Asn Ser Tyr Tyr Ser Trp Tyr His Asp Tyr Gly His Leu Glu Leu 750	2756

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1300 [Street, NW Washington, DC 20005 202,408,4000 fax 202,408,4400 www.finnegan.com ATT CAG CTG CAG CTG GCC ACC CAG TTT GAG AAC TGG TAT AAG AAG 2801 Gln Leu Gln Leu Ala Thr Gln Phe Glu Asn Trp Tyr Lys Lys 760 2846 TAT CAG AAG CCC ATT ATT CAG AGC GAG TAT GGA GCA GAA ACG ATT Tyr Gin Lys Pro Ile Ile Gin Ser Glu Tyr Gly Ala Glu Thr GCA GGG TTT CAC CAG GAT CCA CCT CTG ATG TTC ACT GAA GAG TAC 2891 Ala Gly Phe His Gln Asp Pro Pro Leu Met Phe Thr Glu Glu Tyr 790 800 CAG AAA AGT CTG CTA GAG CAG TAC CAT CTG GGT CTG GAT CAA AAA 2936 Gln Lys Ser Leu Leu Glu Gln Tyr His Leu Gly Leu Asp Gln Lys 810 2981 CGC AGA AAA TAT GTG GTT GGA GAG CTC ATT TGG AAT TTT GCC GAT Arg Arg Lys Tyr Val Val Gly Glu Leu lle Trp Asn Phe Ala Asp 820 830 TTC ATG ACT GAA CAG TCA CCG ACG AGA GTG CTG GGG ATT AAA AAG 3026 Phe Met Thr Glu Gln Ser Pro Thr Arg Val Leu Gly Asn Lys Lys 840 GGG ATC TTC ACT CGG CAG AGA CAA CCA AAA AGT GCA GCG TTC CTT 3071 Phe Thr Arg Gln Arg Gln Pro Lys Ser Ala Ala lle Phe Leu 850 TTG CGA GAG AGA TAC TGG AAG ATT GCC AAT GAA ACC AGG TAT CCC 3116 Leu Arg Glu Arg Tyr Trp Lys lle Ala Asn Glu Thr Arg Tyr Pro 870 CAC TCA GTA GCC AAG TCA CAA TGT TTG GAA AAC AGC CCG TTT ACT 3161 Ser Val Ala Lys Ser Gln Cys Leu Glu Asn Ser Pro Phe Thr 880 890 TGA GCAAGACTGA TACCACCTGC GTGTCCCTTC CTCCCCGAGT CAGGGCGACT 3214 TCCACAGCAG CAGACAAGT GCCTCCTGGA CTGTTCACGG CAGACCAGAA 3264 CGTTTCTGGC CTGGGTTTTG TGGTCATCTA TTCTAGCAGG GAACACTAAA 3314.

- 17. (Withdrawn) A vector containing a nucleic acid as claimed in claim 14.
- 18. (Withdrawn) A host cell containing a nucleic acid as claimed in claim 14 or a vector as claimed in claim 17.
- 19. (Withdrawn) A host cell as claimed in claim 18, which is a BHK, CHO, COS, HeLa, insect, tobacco plant, yeast or *E. coli* cell.
- 20. (Withdrawn) A transgenic mammal with the exception of a human, containing a DNA as claimed in claim 14 or a vector as claimed in claim 17.

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- 21. (Withdrawn) A process for preparing a compound as claimed in claim 1, which comprises
 - a) introducing a nucleic acid as claimed in claim 14 or a vector as claimed in claim 17 into a host cell,
 - b) cultivating the host cell, and
 - c) isolating the compound.
- 22. (Withdrawn) A process for preparing a compound as claimed in claim 1, which comprises
 - a) cultivating a host cell as claimed in claim 18, and
 - b) isolating the compound.
 - 23. (Canceled).
 - 24. (Canceled).
- 25. (Previously Presented) A pharmaceutical composition comprising a compound as claimed in claim 1 and a physiologically acceptable carrier.
- 26. (Previously Presented) A diagnostic aid comprising a compound as claimed in claim 1.
- 27. (Previously Presented) A compound as claimed in claim 6, wherein the lactamase enzyme is a *Bacillys cereus* β -lactamase II.
- 28. (Previously Presented) A compound as claimed in claim 6, wherein the carboxypeptidase enzyme is a carboxypeptidase G2 from *Pseudomonas*.
- 29. (Previously Presented) A compound as claimed in claim 10, which has undergone secretory expression in *Hansenula polymorpha*.

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1300 f Street, NV Washington, DC 20005 202,408,4000 fax 202,408,4400 www.finnegan.com 30. (Previously Presented) A compound as claimed in claim 1, wherein at lease one antigen binding region and at least one prodrug-activating enzyme form an sFv- β -lactamase fusion protein.

31. (Previously Presented) A compound as claimed in claim 11, wherein the chemical glycosylation involves at least one of galactosylation or mannosylation.

32. (Previously Presented) A compound as claimed in claim 12, wherein the chemical glycosylation involves at least one of galactosylation or mannosylation.

33. (Previously Presented) A method of treating cancer comprising administering a compound claimed in claim 1 to a host in need thereof and subsequently administering a prodrug to be activated by the enzyme portion of the compound of claim 1.

34. (Currently Amended) A compound comprising one or more antigen binding regions linked to at least one prodrug-activating enzyme, wherein

- a) the antigen binding regions consist of a single polypeptide chain;
- b) the single polypeptide chain is comprised of a first variable domain, a second variable domain, and a polypeptide linker connecting the first variable domain and the second variable domain; wherein a nucleotide sequence encoding the polypeptide linker is formed by two partially overlapping PCR primers during a PCR reaction that links the first variable domain and the second variable domain; and wherein

c) - a nucleotide sequence encoding the polypeptide linker is formed by two partially overlapping PCR primers during a PCR reaction that

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1300 F Street, NW Washington, DC 20005 202 408 4000 Fax 202 408 4400 www.finnegan.com links the first variable domain and the second variable domain; and wherein

d) c) the compound has a monovalent, bivalent, or multivalent structure.

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